

GenCore version 4.5
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Run on: November 20, 1999, 20:38:59 ; Search time 48.9 Seconds
(without adjustments)

9690.466 Million cell updates/sec

Sequence: 1 gctgacttcctccagcac.....ataaagatactagagaactg 1894

Scoring table: IDENTITY_NUCDX

Searched: 311585 seqs, 125096042 residues

Database : N_Geneseq_36

Pred. NO. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
C 1	78.4	4.1	2567	1	74198	Human ER3 repress
C 2	78.2	4.1	1116	1	V38334	Transcription fac
C 3	78.2	4.1	1907	1	V38335	Transcription fac
C 4	78	4.1	2073	1	O55149	Adenovirus E1a-F
C 5	78	4.1	2064	1	V37087	ElfA1 matrix metal
C 6	76.6	4.0	1904	1	V47071	Human E1a protein
C 7	76.4	4.0	1900	1	O59769	Coding sequence o
C 8	74.2	3.9	2938	1	O50644	Human Hum-F11.1 g
C 9	74.2	3.9	1890	1	O50664	Human Hum-F11.1 g
C 10	73.4	3.9	2410	1	V32688	Polyomavirus PEA3
C 11	68.6	3.6	7422	1	V32689	Mouse ESX transscr
C 12	68.6	3.6	7422	1	V32689	Mouse ESX transscr
C 13	60.6	3.2	852	1	V61378	Human lymphoid-B
C 14	60.4	3.2	2975	1	V15529	Human lymphoid-B
C 15	60.4	3.2	3240	1	V15530	Human lymphoid-B
C 16	60	3.2	2544	1	V47200	Mouse E125 repress
C 17	59.6	3.1	1504	1	V20455	Human c-rets oncog
C 18	53.4	2.8	1447	1	X26551	DNA encoding ch1c2
C 19	53.4	2.8	1538	1	X26552	DNA encoding ch1c2
C 20	49	2.6	1364	1	V20472	DNA encoding ch1c2
C 21	46.2	2.4	1913	1	O15015	Human c-spi oncog
C 22	45.8	2.4	15079	1	O91580	Nuclear factor C/
C 23	45.4	2.4	2386	1	O63477	S. clavicolacti p
C 24	45.4	2.4	2386	1	O63477	Micrococcal p
C 25	45.2	2.4	13904	1	T89415	Streptomyces Yema
C 26	44.2	2.4	13904	1	T89415	Streptomyces Yema
C 27	44.6	2.4	2564	1	V67065	Human alpha1(X)
C 28	43	2.3	16449	1	T67066	Human alpha1(X)
C 29	43	2.3	985	1	V64548	Myobacteriobas
C 30	42.6	2.2	2338	1	O76123	N. tuberculosis
C 31	42.6	2.2	18001	1	O76123	N. clavipes dirgill
C 32	42.6	2.2	2338	1	V23249	HSV L/ST region.
C 33	42.6	2.2	4636	1	X33449	Nephila clavipes
C 34	42	2.2	2004	1	T85356	Oryza sativa L. p
C 35	41.6	2.2	2558	1	T88419	Nephila clavipes
C 36	41.6	2.2	1233	1	V00246	Human g1a1 cell
C 37	41.6	2.2	1233	1	V00246	Human g1a1 cell
C 38	41.6	2.2	1937	1	V09317	Human Ret ligand
C 39	41.6	2.2	1937	1	V09317	Human Ret ligand
C 40	41.4	2.2	1939	1	V09315	gDNFR-alpha cDNA
C 41	41.4	2.2	53789	1	V21317	gDNFR-alpha cDNA
C 42	41.2	2.2	738	1	X39714	Amlycotic cancer
C 43	41.2	2.2	937	1	X39716	Amlycotic cancer
C 44	41.2	2.2	1985	1	O37460	Genetic cancer as

[illegible]

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RESULT      2
V8354      ID V8354 standard; cDNA: 1116 Bp.
DE 09-NOV-1998 (first entry)
DE Transcription factor ESX gene open reading frame
KW ESX: epithelial-restricted with serine box; transcription factor;
KW ESX: human; epithelial cancer; breast cancer; adenocarcinoma;
KW diagnosis; therapy: ss.
OS Homo sapiens.
PI Benz CC: Chag9 C: Scott GK;
PI WPI: 98-322755/28.
PI P-PSDB: W60677-79.
DE Nucleic acid encoding the ESX transcription activator -
DE over-expressed in epithelial, especially breast, cancers, also
DE related polypeptide(s), antibodies, vectors and transformed cells,
DE useful for diagnosis and treatment of cancer
DE Title: ESX 1; 120pp; English.
DE Summary: This is a full length cDNA of the newly isolated ESX gene that
DE codes for a novel transcription factor (see W60677) associated

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CC With the etiology of cancers, including epithelial cancers.
CC The gene is a member of the ESX gene family designated ESX for
CC epithelial-restricted with serine box. The gene is located on
CC chromosome 19p32, a region amplified in 50% of early breast cancers.
CC ESX is an early indicator of breast cancer, being expressed at the
CC ductal carcinoma in situ stage. Dysfunction of the ESX gene that
CC results in increased expression is indicative of epithelial cancer
CC especially breast cancer but also gastric, ovarian and lung
CC cancers. The gene is also a prognostic factor for the ESX open
CC reading frame was deduced from a cDNA library. The ESX gene
CC origin that was identified in an expressed sequence tag database
CC search using human Ets-2, and from a clone obtained by 5' RACE of
CC placental polyA mRNA. Antibodies, antisense nucleic acids or
CC inactive ESX mutant proteins that inhibit activity of ESX can be
CC used to inhibit growth and proliferation of neoplastic cells.
CC Antibodies are used to detect ESX polypeptides, particularly for diagnosis
CC and monitoring of cancer. Polynucleotides are used for in vitro
CC detect gross abnormalities in chromosome 19p32 for in vitro
CC screening for predisposition to cancer, and to generate ESX protein
CC in vivo from gene therapy vectors. ESX protein, genes and cDNA
CC can also be used to screen for specific modulators, especially
CC potential anticancer agents.
CC Sequence 1116 Bp; 235 A; 322 C; 335 G; 204 T;
SQ
Query Match 4.1%; Score 78.2; DB 1; Length 1116;
Best Local Similarity 59.8%; Pred. No. 1.2e-07;
Matches 150; Conservative 0; Mismatches 98; Indels 3; Gaps 1;
QY 1159 catccacactggtggcagttctctcctcagagagtgctactcaagcccccacagctagtcgcgtt 1218
Db 816 CACCCACCTGTGGAGATTCCTCCGGAACATCTCCACCCGACTCTCAAGAGGCTT 875
QY 1219 cattagtggtctacacaggaagagcattcaaataggactcaagcaggtggc 1278
Db 876 CATGAATGGGAGAAATCGCATGAGAGGCTTCCTCAACTCTCCGCTCCGAGGCTGTGGC 935
QY 1279 ccgagctgtagggcattccgcagacgcgtccgcgtctgaactagcaagctagtcgcgtc 1338
Db 936 CCAACTATGGGGCCCAAAAGAAAGAACATCACTATGCAAGAGCTGAGCGCGGC 995
QY 1339 catccgcagagattacaaagagcattccatccgcagacacattcccaagcgtcgt 1398
Db 996 CATGAG--GTCATCTACAAACGAGGAATCTGGACAGGGTGGATGAGCCGCGACATCTGT 1052
QY 1399 ctaccagcttcg 1409
Db 1053 CTACAGATTG 1063
DE 09-NOV-1998 (first entry)
DE Transcription factor ESX cDNA (ORF and untranslated regions).
DE ESX: epithelial-restricted with serine box; transcription factor;
DE ESX: human; epithelial cancer; breast cancer; adenocarcinoma;
DE diagnosis; therapy: ss.
OS Homo sapiens.
PI Benz CC: Chag9 C: Scott GK;
PI WPI: 98-322755/28.
PI P-PSDB: W60677-79.
DE Nucleic acid encoding the ESX transcription activator -
DE over-expressed in epithelial, especially breast, cancers, also
DE related polypeptide(s), antibodies, vectors and transformed cells,
DE useful for diagnosis and treatment of cancer
DE Title: ESX 1; 120pp; English.
DE Summary: This is a full length cDNA of the newly isolated ESX gene that
DE codes for a novel transcription factor (see W60677) associated

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OY 1370 c9gang 1375 .
DB 1306 CAGAG 1311

RESULT 8
ID 050644 standard: cDNA: 2938 BP.
AC 050644:
DE 26-MAY-1994 (first entry)
KW Human Hum-Flil-1 gene clone BM025.
KW chromosomal translocation: chimeric; Ewing sarcoma;
KW EWS gene: malignant melanoma; hum-flil-1;
KW primitive peripheral neuroectodermal tumour; human chromosome 11;
OS Homo sapiens.
FH key Location/Qualifiers
FT cds 143..1501
FT /*cag- a
FT /*product= HUM-FLI-1
FT /*cag- b
FT /*cag- d
FT poly-a_signal
FT 1908..1913
FT W09323549-A.
FT 25-NOV-1993.
FT 19-MAY-1993: FR-006123.
FT 20-MAY-1993: FR-006123.
FT (CNRS ) CNRS CENT NAT RECH SCI.
FT Aurias A., Delattre O., Desmaza C., Melot T., Peter M;
FT Plogastrel B., Thomas G., Zucman J;
FT MPI: 93-386580/48.
FT New nucleic acid of EWS gene and its hybrid(s) - contg. gene
FT sequence involved in chromosomal translocation, also derived
FT from RNA, probes, fusion proteins etc., for diagnosis and treatment
FT of Ewing sarcoma and melanoma
FT Discloure: Fig 7; 123pp: French.
FT The probe 11R1 was used to screen a human marrow cDNA library
FT (Clontech cat. # HL1058). The clone BM025 was identified and
FT sequenced. It represents the entire coding region together with
FT and 3' UTR of the Hum-Flil-1 gene.
FT Sequence 2938 BP: 847 A; 692 C; 675 G; 724 T;
SO

Query Match 3.9%; Score 74.2; DB 1; Length 2938;
Best Local Similarity 62.9%; Pred. No. 9e-07;
Matches 134; Conservative 0; Mismatches 73; Indels 6; Gaps 1;

OY 1156 gccctccacccctggtcgtctctcaaggaggtgtctactcaagcccccagcgtatggcgcg 1215
DB 979 ctcacatctacacgtgtgacatctctctcgcagacgtctccgcacagccgcacgcacgtg 1038
OY 1216 ctctacatctacgtgtgctcaacagagagaggtcgtctctcaaaattgaggtcagcccggt 1275
DB 1039 TATCACTGCGAGGAGGCCAAGCCAGCGGAG-----TTCAAAATGACGACCCCGATGAGGT 1092
OY 1276 ggcgcgcgtgtgtgggcatcgcagaaacgcgtccgcacatgactcagacagagcgtgagcgcg 1335
DB 1093 GCGCAGCGCGTGTGGCGACCGGAAAGCAAGCCCAACATGATATACGACACTGAGCGG 1152
OY 1336 ctctctcgcgcgcgtatctcagaagaggtcattc 1368
DB 1153 GCGCTCTCGTATTAATGATATGATAAACATTAAT 1185

RESULT 9
ID 050662 standard: DNA: 1890 BP.
AC 050662:
DE 26-MAY-1994 (first entry)
KW Human Hum-Flil-1 gene clone BM025.
KW chromosomal translocation: chimeric; Ewing sarcoma;
KW EWS gene: malignant melanoma; hum-flil-1;
OS Homo sapiens.
FH key Location/Qualifiers
FT cds 143..1501
FT /*cag- a
FT /*product= HUM-FLI-1
FT /*cag- b
FT /*cag- d
FT poly-a_signal
FT 1908..1913
FT W09323549-A.
FT 25-NOV-1993.
FT 19-MAY-1993: FR-006123.
FT 20-MAY-1993: FR-006123.
FT (CNRS ) CNRS CENT NAT RECH SCI.
FT Aurias A., Delattre O., Desmaza C., Melot T., Peter M;
FT Plogastrel B., Thomas G., Zucman J;
FT MPI: 93-386580/48.
FT New nucleic acid of EWS gene and its hybrid(s) - contg. gene
FT sequence involved in chromosomal translocation, also derived
FT from RNA, probes, fusion proteins etc., for diagnosis and treatment
FT of Ewing sarcoma and melanoma
FT Discloure: Fig 7 and Fig 13; 123pp: French.
FT The probe 11R1 was used to screen a human marrow cDNA library
FT (Clontech cat. # HL1058). The clone BM025 was identified and
FT sequenced. It represents the entire coding region together with
FT and 3' UTR of the Hum-Flil-1 gene.
FT Sequence 2938 BP: 847 A; 692 C; 675 G; 724 T;
SO

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KW primitive peripheral neuroectodermal tumour; human chromosome 11;
OS Homo sapiens.
FH key Location/Qualifiers
FT exon 1..39
FT /*cag- a
FT /*product= HUM-FLI-1
FT /*cag- b
FT /*cag- d
FT poly-a_signal
FT 1908..1913
FT W09323549-A.
FT 25-NOV-1993.
FT 19-MAY-1993: FR-006123.
FT 20-MAY-1993: FR-006123.
FT (CNRS ) CNRS CENT NAT RECH SCI.
FT Aurias A., Delattre O., Desmaza C., Melot T., Peter M;
FT Plogastrel B., Thomas G., Zucman J;
FT MPI: 93-386580/48.
FT New nucleic acid of EWS gene and its hybrid(s) - contg. gene
FT sequence involved in chromosomal translocation, also derived
FT from RNA, probes, fusion proteins etc., for diagnosis and treatment
FT of Ewing sarcoma and melanoma
FT Discloure: Fig 7 and Fig 13; 123pp: French.
FT The probe 11R1 was used to screen a human marrow cDNA library
FT (Clontech cat. # HL1058). The clone BM025 was identified and
FT sequenced. It represents the entire coding region together with
FT and 3' UTR of the Hum-Flil-1 gene.
FT Sequence 2938 BP: 847 A; 692 C; 675 G; 724 T;
SO

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CC exons from the two genes, as happens after specific chromosomal
CC translocations, can be predicted (see 950611-950678). The sequences
CC at fusion junctions of other observed translocations are given in
CC 950679-950683). NOTE: The 1890 residue sequence 950665 does not
CC appear in the specification; it is a combination of the Hum-Flt-1
CC cDNA sequence (Fig 9, 950644) and the individual intron sequences
CC with their intron-exon junctions (Fig 13). 458 G; 414 T;
50 Sequence 1890 bp: 437 A; 451 C; 456 G; 457 A; 458 G; 459 G; 460 G; 461 G; 462 G; 463 G; 464 G; 465 G; 466 G; 467 G; 468 G; 469 G; 470 G; 471 G; 472 G; 473 G; 474 G; 475 G; 476 G; 477 G; 478 G; 479 G; 480 G; 481 G; 482 G; 483 G; 484 G; 485 G; 486 G; 487 G; 488 G; 489 G; 490 G; 491 G; 492 G; 493 G; 494 G; 495 G; 496 G; 497 G; 498 G; 499 G; 500 G; 501 G; 502 G; 503 G; 504 G; 505 G; 506 G; 507 G; 508 G; 509 G; 510 G; 511 G; 512 G; 513 G; 514 G; 515 G; 516 G; 517 G; 518 G; 519 G; 520 G; 521 G; 522 G; 523 G; 524 G; 525 G; 526 G; 527 G; 528 G; 529 G; 530 G; 531 G; 532 G; 533 G; 534 G; 535 G; 536 G; 537 G; 538 G; 539 G; 540 G; 541 G; 542 G; 543 G; 544 G; 545 G; 546 G; 547 G; 548 G; 549 G; 550 G; 551 G; 552 G; 553 G; 554 G; 555 G; 556 G; 557 G; 558 G; 559 G; 560 G; 561 G; 562 G; 563 G; 564 G; 565 G; 566 G; 567 G; 568 G; 569 G; 570 G; 571 G; 572 G; 573 G; 574 G; 575 G; 576 G; 577 G; 578 G; 579 G; 580 G; 581 G; 582 G; 583 G; 584 G; 585 G; 586 G; 587 G; 588 G; 589 G; 590 G; 591 G; 592 G; 593 G; 594 G; 595 G; 596 G; 597 G; 598 G; 599 G; 600 G; 601 G; 602 G; 603 G; 604 G; 605 G; 606 G; 607 G; 608 G; 609 G; 610 G; 611 G; 612 G; 613 G; 614 G; 615 G; 616 G; 617 G; 618 G; 619 G; 620 G; 621 G; 622 G; 623 G; 624 G; 625 G; 626 G; 627 G; 628 G; 629 G; 630 G; 631 G; 632 G; 633 G; 634 G; 635 G; 636 G; 637 G; 638 G; 639 G; 640 G; 641 G; 642 G; 643 G; 644 G; 645 G; 646 G; 647 G; 648 G; 649 G; 650 G; 651 G; 652 G; 653 G; 654 G; 655 G; 656 G; 657 G; 658 G; 659 G; 660 G; 661 G; 662 G; 663 G; 664 G; 665 G; 666 G; 667 G; 668 G; 669 G; 670 G; 671 G; 672 G; 673 G; 674 G; 675 G; 676 G; 677 G; 678 G; 679 G; 680 G; 681 G; 682 G; 683 G; 684 G; 685 G; 686 G; 687 G; 688 G; 689 G; 690 G; 691 G; 692 G; 693 G; 694 G; 695 G; 696 G; 697 G; 698 G; 699 G; 700 G; 701 G; 702 G; 703 G; 704 G; 705 G; 706 G; 707 G; 708 G; 709 G; 710 G; 711 G; 712 G; 713 G; 714 G; 715 G; 716 G; 717 G; 718 G; 719 G; 720 G; 721 G; 722 G; 723 G; 724 G; 725 G; 726 G; 727 G; 728 G; 729 G; 730 G; 731 G; 732 G; 733 G; 734 G; 735 G; 736 G; 737 G; 738 G; 739 G; 740 G; 741 G; 742 G; 743 G; 744 G; 745 G; 746 G; 747 G; 748 G; 749 G; 750 G; 751 G; 752 G; 753 G; 754 G; 755 G; 756 G; 757 G; 758 G; 759 G; 760 G; 761 G; 762 G; 763 G; 764 G; 765 G; 766 G; 767 G; 768 G; 769 G; 770 G; 771 G; 772 G; 773 G; 774 G; 775 G; 776 G; 777 G; 778 G; 779 G; 780 G; 781 G; 782 G; 783 G; 784 G; 785 G; 786 G; 787 G; 788 G; 789 G; 790 G; 791 G; 792 G; 793 G; 794 G; 795 G; 796 G; 797 G; 798 G; 799 G; 800 G; 801 G; 802 G; 803 G; 804 G; 805 G; 806 G; 807 G; 808 G; 809 G; 810 G; 811 G; 812 G; 813 G; 814 G; 815 G; 816 G; 817 G; 818 G; 819 G; 820 G; 821 G; 822 G; 823 G; 824 G; 825 G; 826 G; 827 G; 828 G; 829 G; 830 G; 831 G; 832 G; 833 G; 834 G; 835 G; 836 G; 837 G; 838 G; 839 G; 840 G; 841 G; 842 G; 843 G; 844 G; 845 G; 846 G; 847 G; 848 G; 849 G; 850 G; 851 G; 852 G; 853 G; 854 G; 855 G; 856 G; 857 G; 858 G; 859 G; 860 G; 861 G; 862 G; 863 G; 864 G; 865 G; 866 G; 867 G; 868 G; 869 G; 870 G; 871 G; 872 G; 873 G; 874 G; 875 G; 876 G; 877 G; 878 G; 879 G; 880 G; 881 G; 882 G; 883 G; 884 G; 885 G; 886 G; 887 G; 888 G; 889 G; 890 G; 891 G; 892 G; 893 G; 894 G; 895 G; 896 G; 897 G; 898 G; 899 G; 900 G; 901 G; 902 G; 903 G; 904 G; 905 G; 906 G; 907 G; 908 G; 909 G; 910 G; 911 G; 912 G; 913 G; 914 G; 915 G; 916 G; 917 G; 918 G; 919 G; 920 G; 921 G; 922 G; 923 G; 924 G; 925 G; 926 G; 927 G; 928 G; 929 G; 930 G; 931 G; 932 G; 933 G; 934 G; 935 G; 936 G; 937 G; 938 G; 939 G; 940 G; 941 G; 942 G; 943 G; 944 G; 945 G; 946 G; 947 G; 948 G; 949 G; 950 G; 951 G; 952 G; 953 G; 954 G; 955 G; 956 G; 957 G; 958 G; 959 G; 960 G; 961 G; 962 G; 963 G; 964 G; 965 G; 966 G; 967 G; 968 G; 969 G; 970 G; 971 G; 972 G; 973 G; 974 G; 975 G; 976 G; 977 G; 978 G; 979 G; 980 G; 981 G; 982 G; 983 G; 984 G; 985 G; 986 G; 987 G; 988 G; 989 G; 990 G; 991 G; 992 G; 993 G; 994 G; 995 G; 996 G; 997 G; 998 G; 999 G; 1000 G; 1001 G; 1002 G; 1003 G; 1004 G; 1005 G; 1006 G; 1007 G; 1008 G; 1009 G; 1010 G; 1011 G; 1012 G; 1013 G; 1014 G; 1015 G; 1016 G; 1017 G; 1018 G; 1019 G; 1020 G; 1021 G; 1022 G; 1023 G; 1024 G; 1025 G; 1026 G; 1027 G; 1028 G; 1029 G; 1030 G; 1031 G; 1032 G; 1033 G; 1034 G; 1035 G; 1036 G; 1037 G; 1038 G; 1039 G; 1040 G; 1041 G; 1042 G; 1043 G; 1044 G; 1045 G; 1046 G; 1047 G; 1048 G; 1049 G; 1050 G; 1051 G; 1052 G; 1053 G; 1054 G; 1055 G; 1056 G; 1057 G; 1058 G; 1059 G; 1060 G; 1061 G; 1062 G; 1063 G; 1064 G; 1065 G; 1066 G; 1067 G; 1068 G; 1069 G; 1070 G; 1071 G; 1072 G; 1073 G; 1074 G; 1075 G; 1076 G; 1077 G; 1078 G; 1079 G; 1080 G; 1081 G; 1082 G; 1083 G; 1084 G; 1085 G; 1086 G; 1087 G; 1088 G; 1089 G; 1090 G;

Query Match	Score 74.2	DB 1	Length 1890
Best Local Similarity	62.9%	Prod. No. 8.5e-07	
Matches 136	Conservative	0	Mismatches 73; Indels 6; Gaps 1
QY	1156 gccacacacccctgtggagcttcttcacaggatttgctactaacccccacgatagtgcg	1215	
DB	1338 gccatctccctgtgtccagcaatctcttcacagctctctcagcaagccgacacgacgacgtg	1397	
QY	1216 cttatctatgtgtccctggacacatggagacatctcaataattcaggacacgacccagc	1275	
DB	1398 tttctactctggagacggacacacacggcgac-----ttcattatctatgacacccctgac	1451	
QY	1276 gggccggagctgtgggacatccgcagaagacgtgccgcgataaacctagcaagaactgtggcgc	1335	
DB	1452 gccctacgcgcctggcgscagcgaggaagcaaaccccaactcattatgacacagctcgagccg	1511	
QY	1336 ctccatctgcgcagattctacaagaagagcatcat	1368	
DB	1512 gccctctgcgttatctatctatgataataaataacattat	1544	

RESULT	10	
V32688		
ID	V32688 standard; cDNA; 2410 BP.	
AC	V32688	
DT	20-OCT-1998 (first entry)	
DR		
PT	Polyomavirus PEA3 cDNA.	
PR	Polyomavirus PEA3 enhancer activator. PEA3; tumour; suppressor; inhibitor;	
PM	polyomavirus. HER-2; neu promoter; metastasis; cancer; ss.	
KS	Polyomavirus16	
OW		
FM	Key	Location/Qualifiers
FT	CDS	126..-1793
FT		..cag- a
FT		/product= PEA3
FN		/note= "polyomavirus enhancer activator"
PD	NC09830585-A2.	
PD	15-JUL-1998.	
PR	15-JAN-1997; US928085.	
PR	10-JAN-1997; US928085.	
PA	(TEPA3) UNITV TEPA3 SYSTEM.	
PI	Hung M, Xing X.	
PI	WPI: 98-399061/34.	
DR	P-PDB: M49010.	
DR		
PT	Method for representing transformation of cells - by contacting cell	
PT	with polyoma-virus enhancer activator 3, useful for: e.g. treating	
PT	preventing cancer, tumorigenesis and metastasis	
PT	disclosure: Page 69-70: 3Bp; English.	
PT	This sequence encodes a polyomavirus enhancer activator, PEA3. This	
PT	sequence is used in a method for representing transformation in a cell	
PT	which involves contacting the cell with PEA3 to inhibit a transformed	
PT	cell. This sequence can also be used in a method to suppress the	
PT	growth of a tumour comprising introducing to the tumour, a	
PT	polyomavirus encoding nucleic acid having the expression of PEA3 in the mammal,	
PT	results in a decrease in the growth of the tumour. PEA3 can regulate	
PT	the HER-2/neu promoter by transcriptional repression. PEA3 as a tumour	
PT	repressor. The PEA3 can be used for reducing a transforming	
PT	tumorigenic or metastatic potential of a cell. It can be used for the	
PT	prevention and treatment of cancer.	
PT	tumorigenesis and metastasis.	
Sequence	2410 BP: 521 A:	744 C: 645 G: 500 T:

Query Match

Best Local Similarity	59.9%	Pred. No. 1,36-06
Matches 160:	Conservative 0:	Mismatches 101: Indels 6: Gaps 2:
Oy 1109	gagagagagctgagacagacagagatgagatcatcatgctctcgagagccatccactcgtg 1168	
Db 1308	GAAGGATGTGAGCTTTCTCGGAGAGGCGCCACCTTACACAGCGCCGAGGTCTCTTACACTG 1367	
Oy 1189	tggcagttcccaagggtgtgtcatctctcagccccaagacagatgagcgcctcatatgtgtg 1228	
Dy 1368	TGGCAGTTTCT--GGTGGCCCTCTGTGATACCCACAAAGCTCATTTCAATTCGCTTGG 1424	
Oy 1229	ctcaacacagagagagagatctcaaatctgaagcatcagacacagagtgagccagctgtgtg 1288	
Dy 1455	ACAGGCGCGGAGATGG--AGTTTAACTATATGAACTGTAAAGAGTGTCCAGGCTCTGG 1481	
Oy 1289	ggcatccgcagagacgcgtccgcgcatacgaactgaacgaagagctgacgcgtcatccgcag 1348	
Dy 1482	GGATTCGAGAGAGACGGCCACCATGAATATGACAAAGTACGACCGCTCCGTCCGATAC 1541	
Oy 1349	tattacagagagagcatctccagag 1375	
Dy 1542	TATTATGAAAGCTCATTCAGAG 1568	
RESULT 11		
11	11	
AC	V38365 standard: cDNA; 7752 BP.	
AC	V38365:	
DE	09-NOV-1998 (first entry)	
DE	Mouse ESX transcription factor gene.	
KM	ESX: epithelial-restricted with serine box; transcription factor;	
KM	ERTS: mouse; epithelial cancer; breast cancer; adenocarcinoma;	
OM	diagnosis: therapy: ss.	
PH	Key SP.	
PH	location/Qualifiers	
FT	promoter	2553..2889
FT	/*tag= a	3604..7115
FT	CD5	
FT	/*tag= b	3200..3035
FT	/*note= "contains introns"	
FT	exon	3036..3606
FT	/*tag= c	
FT	intron	3607..4148
FT	/*tag= d	
FT	exon	4149..4503
FT	/*tag= e	
FT	intron	4504..4594
FT	/*tag= f	
FT	exon	4595..4782
FT	/*tag= g	
FT	intron	4783..4905
FT	/*tag= h	
FT	exon	4906..5052
FT	/*tag= i	
FT	intron	5053..5142
FT	/*tag= j	
FT	exon	5143..5168
FT	/*tag= k	
FT	intron	5169..5214
FT	/*tag= l	
FT	exon	5215..5260
FT	/*tag= m	
FT	intron	5261..5316
FT	/*tag= n	
FT	exon	5317..5362
FT	/*tag= o	
FT	intron	5363..5418
FT	/*tag= p	
FT	exon	5419..5464
FT	/*tag= q	
FT	intron	5465..5520
FT	/*tag= r	
FT	exon	5521..5576
FT	/*tag= s	
FT	intron	5577..5632
FT	/*tag= t	
FT	exon	5633..5688
FT	/*tag= u	
FT	intron	5689..5744
FT	/*tag= v	
FT	exon	5745..5800
FT	/*tag= w	
FT	intron	5801..5856
FT	/*tag= x	
FT	exon	5857..5912
FT	/*tag= y	
FT	intron	5913..5968
FT	/*tag= z	
FT	exon	5969..6024
FT	/*tag= AA	
FT	intron	6025..6080
FT	/*tag= AB	
FT	exon	6081..6136
FT	/*tag= AC	
FT	intron	6137..6192
FT	/*tag= AD	
FT	exon	6193..6248
FT	/*tag= AE	
FT	intron	6249..6304
FT	/*tag= AF	
FT	exon	6305..6360
FT	/*tag= AG	
FT	intron	6361..6416
FT	/*tag= AH	
FT	exon	6417..6472
FT	/*tag= AI	
FT	intron	6473..6528
FT	/*tag= AJ	
FT	exon	6529..6584
FT	/*tag= AK	
FT	intron	6585..6640
FT	/*tag= AL	
FT	exon	6641..6696
FT	/*tag= AM	
FT	intron	6697..6752
FT	/*tag= AN	
FT	exon	6753..6808
FT	/*tag= AO	
FT	intron	6809..6864
FT	/*tag= AP	
FT	exon	6865..6920
FT	/*tag= AQ	
FT	intron	6921..6976
FT	/*tag= AR	
FT	exon	6977..7032
FT	/*tag= AS	

us-09-126-945-1.rng

CC	adenocarcinoma) and of unfavourable prognosis.
CC	The ESX open
CC	reading frame was deduced from a cDNA clone of foetal liver-lymph
CC	origin that was identified in an expressed sequence tag database
CC	search using human Ets-2, and from a clone obtained by 5'RACE of
CC	placental polyA mRNA. Antibodies, antisense nucleic acids or
CC	inactive ESX mutant proteins that inhibit activity of ESX can be
CC	used to inhibit growth and proliferation of neoplastic cells,
CC	antibodies were used to examine ESX polypeptides, polynucleotides and
CC	antibodies were used to examine ESX polypeptides, polynucleotides and
CC	antibodies were used to examine ESX polypeptides, polynucleotides and
CC	monitors of cancer. Polynucleotides may also be used to detect
CC	gross abnormalities in chromosome 11, e.g. for in vitro
CC	screening for predisposition to cancer, and to generate ESX protein
CC	in vivo from gene therapy vectors. ESX protein, genes and cDNA
CC	can also be used to screen for specific modulators, especially
CC	potential anticancer agents.
CC	Sequence 1116 bp; 235 A; 322 C; 335 G; 204 T;
SQ	
Query Match	4.1%; Score 78.2; DB 1; Length 1116;
Best Local Similarity	59.8%; Pred.No.1.2e-07;
Matches 150:	Conservative 0; Mismatches 98; Indels 3; Gaps 1;
OY	1159 catccactcgtgcagcttctccaaggaaatgcatcccaagcccaccgaacgacctc 1218
Dd	816 CACCAACCTGTGGAAATTCATTCGCGCATCTCAACACCGAGACTCAACAAGGCT 875
OY	1219 cattcagtgcctacaagaaggaagggatccttcataacttttggaactcaaccaa 1278
Dd	876 CATGAAGTGGGGAATGCATGAGAAGCGGCTTTCTTAATTCTCGCGCTCCAGAGCTGTGGC 935
OY	1279 ccagcgctgtggggcctccgacagaaacgcgtccgcgcgaactacgaacagctgagccgc 1338
Dd	936 CCACATGATGGGCCAAAAGAAAAAGAACACACATCACTTCGAAACATCTAAGC 995
OY	1339 catccgcgcagatattacaagaagaggcctcctcctcgcgaagcagaactccgccgcctc 1398
Dd	996 CATGAG--GACTCATCTCAACAAAGGAGATCTCGAAGCGTGATGCGCGCACTCT 1052
OY	1399 ctaccacgctc 1409
Dd	1053 CTACAGTTG 1063
RESULT	
V335	3
ID	V335 standard; cDNA, 1907 bp.
AC	V3355.
DT	09-NOV-1998 (first entry)
KD	Transcription factor, ESX cDNA (ORF and untranslated regions);
DE	ESX; epithelial-restricted with serine box; transcription factor;
KM	ESX; human; epithelial cancer; breast cancer; adenocarcinoma;
KW	diagnosis; Therapy; ss.
FT	Key words: Esx proteins.
FT	5'UTR Location/Qualifiers
FT	1..95 /tag-a
FT	CDS 96..1206 /tag-b
FT	/note= "Cabin 3"
FT	1207..1907 /tag-c
FT	3'UTR
PD	MO9823187-A2.
PD	04-JUN-1998.
PF	26-NOV-1997; U21865.
PR	25-NOV-1997; US-031504.
PR	27-NOV-1996; US-031504.
PT	(RESC) UNIV CALIFORNIA,
PI	Benz CG, Chang C, Scott GK,
PI	Ben J, et al. 1997/28.
PIDB	P-IDB: M60672
DR	Nucleic acid encoding the ESX transcription activator -
DT	related polypeptide(s), antibodies, vectors and transformed cells.

	/note-	*Claimed sequence
FT	J05328975-A.	
PN	14-DEC-1993.	
PD	02-JUN-1992.	165453.
PF	02-JUN-1992.	JP-165453.
PR		

CC The present sequence encodes the Elaf protein, which is a matrix
CC metalloproteinase regulator, the infiltration and metastasis of
CC expression products of the Elaf gene. This may be accomplished by
CC inducing antisense DNA or RNA for the Elaf gene, a decoy gene
CC expressing the DNA binding region of the Elaf protein, the target
CC DNA for the DNA binding region of the Elaf protein or ribosomes
CC corresponding to the Elaf gene mRNA. Cancer can be diagnosed by
CC detecting the expression of the Elaf gene in tumor cells.
CC These methods may be used in the treatment and diagnosis of cancer
CC e.g. mammary cancer, fibrosarcoma, osteosarcoma, lung cancer, etc.
CC Sequence: 2064 bp; 437 A; 648 C; 569 G; 420 T.

Query Match	4.1%	Score 78:	DB 1:	Length 2064:
Best Local Similarity	76.2%	Pred. No. 1.5e-07:		
Matches 96:	Conservative 0:	Mismatches 30:	Indels 0:	Gaps 0:

FT /number=7
 FT exon 6256..6450
 FT //tag=9
 FT /number=8
 FT intron 6451..6998
 FT //tag=1
 FT /number=8
 FT exon 6999..7155
 FT //tag=8
 FT /number=9
 FT M0823782.A2.
 PD 04-JUN-1998.
 PD 26-NOV-1997; U21865.
 PR 25-NOV-1997; US-031504.
 PR 27-NOV-1996; US-031504.
 PR RECC (J) UNIV CALIFORNIA.
 PR 58-10-1997; US-031504.
 DR WPI: 98-12275/28.
 DR P-F508: w60880.
 PT Nucleic acid encoding the ESX transcription activator -
 PT over-expressed in epithelial, especially breast, cancers, also
 PT related polypeptide(s), antibodies, vectors and transformed cells,
 PT useful for diagnosis and treatment of cancer
 FT
 FT Claim 18: Page 85-90; 140pp; English.
 CC 9 exon that specify a transcript of about 4.9 kb of DNA incorporating at least
 CC encoding the 371-amino acid mESX protein (see w60880). The gene
 CC is a member of the ETS family and is designated Etx for
 CC epithelial-restricted with serine box. In humans, a dysfunction of
 CC the Etx gene (see V18354) that results in increased expression is
 CC indicative of epithelial cancer (especially breast cancer but
 CC also gastric, ovarian and lung adenocarcinoma) and of unfavorable
 CC prognosis. The gene encodes a protein that inhibits cell growth and
 CC tumor proteins that inhibit activity of human ESX can be used to
 CC inhibit growth and proliferation of neoplastic cells, particularly
 CC cancer. ESX polypeptides, polynucleotides and antibodies are also
 CC used to detect ESX, particularly for diagnosis and monitoring of
 CC cancer. They can also be used to screen for specific modulators,
 CC especially potential anticancer agents. Transgenic animals are
 CC also provided.
 SO Sequence 7752 BP; 1768 A; 2050 C; 2019 G; 1914 T;

Query Match 3.6%; Score 68.6; DB 1; Length 7752;
 Best Local Similarity 61.5%; Pred. No. 1.4e-05;
 Matches 110; Conservative 0; Mismatches 69; Indels 0; Gaps 0;
 Oy 1163 caccctgagcagttctcacaaggatgttactactaccccccacagctatggccgttcatt 1222
 Db 6271 CACTGTGCGACATTTATCCGACACATCTTCAATGACCCGACGCTACAGAGAGCCTCATG 6330
 Oy 1223 aggtgtctcaacaaagaaagagcctctcacaattgagctcagccagccagccg 1262
 Db 6331 AAGTGGAGACACCGACAGAGGCTGTCTTCAAGTTCTTCCCTCAGAGCGCTGCCCA 6390
 Oy 1263 ctgtgtgagcctcccaagaacgctcccgcaagctcagcagcctgagccgctcat 1341
 Db 6391 CTTCTGGCCGCAAGAGAGAACAGACAGCATGACTTATGAGAGCTGACGCGACAT 6449
 RESULT 12
 V58521/c
 ID V58521 standard: cDNA: 852 BP.
 AC V58521:
 DT 08-DEC-1998 (first entry)
 DB prostate tumour specific gene clone p18.
 KW prostate tumour specific gene; human; prostate cancer; detection;
 therapy; ss.
 OS Homo sapiens.
 PD 27-AUG-1998.
 PR 25-FEB-1998; U03690.

PR 09-FEB-1998; US-904809.
 PR 25-FEB-1997; US-806596.
 PR 01-AUG-1997; US-904809.
 PA (CORI-) CORIXA CORP.
 PI Dillon DC, Xu J.
 PI WPI: 98-180805/41.
 DR Novel human prostate specific tumour protein and fragments - useful
 PT for diagnosis and treatment of prostate cancer
 PS Claim 1: Page 55; 141pp; English
 CC This sequence represents a human prostate tumour specific gene, and can
 CC be used in the method of the invention. The method is for detecting
 CC prostate cancer comprises contacting a biological sample with an agent
 CC able to bind an immunogenic portion of a prostate protein (such as
 CC encoded by this sequence). An antibody which binds to an immunogen-
 CC portion of the prostate protein, and the method can be used to detect,
 CC monitor progression of, or treat, prostate cancers. The antibody may
 CC be conjugated to a therapeutic agent for use in therapy of prostate
 CC cancers
 SO Sequence 852 BP; 174 A; 195 C; 206 G; 274 T;

Query Match 3.2%; Score 60.6; DB 1; Length 852;
 Best Local Similarity 56.2%; Pred. No. 0.00043;
 Matches 114; Conservative 0; Mismatches 89; Indels 0; Gaps 0;
 Oy 1163 caccctgagcagttctcacaaggatgttactactaccccccacagctatggccgttcatt 1222
 Db 388 CACTTATGCAATTCATCCGACATCTCTTCAACCCACAGACAGACCGAGATTAA 329
 Oy 1223 aggtgtctcaacaaagaaagagcctctcacaattgagctcagccagctgagcccg 1262
 Db 328 AAGTGGAGACACGATCTGAGAGGCTGTCTTCAAGTTCAAGAGAGCCTCATG 269
 Oy 1263 ctgtgtgagcctcccaagaacgctcccgcaagctcagcagcctgagccgctcat 1342
 Db 268 CTTATGGGATTAAGAGAGAACAGACAGCATGACTTATGAGAGCTGACGCGACTATG 209
 Oy 1343 cgcagatctacaaagagagcat 1365
 Db 208 AGATTACTACTACAAAGAGAAAT 186

RESULT 13
 V61167/c
 ID V61167 standard: cDNA: 852 BP.
 AC V61167:
 DT 06-JAN-1999 (first entry)
 DB cDNA sequence of prostate tumour clone p18.
 KW prostate; cancer; tumour; vaccine; immunogen; clone; ss.
 OS Homo sapiens.
 PD W08217093.A2.
 PR 25-FEB-1998; U03492.
 PR 09-FEB-1998; US-020956.
 PR 25-FEB-1997; US-806099.
 PR 01-AUG-1997; US-904804.
 PA (CORI-) CORIXA CORP.
 PI Dillon DC, Xu J.
 PI WPI: 98-609886/51.
 DR Used in a vaccine for the treatment of prostate cancer
 PT polypeptides comprising immunogenic portions of prostate proteins -
 PT used in a vaccine for the treatment of prostate cancer
 PS Claim 1: Page 13; 120pp; English
 CC The present sequence is a new DNA which encodes an immunogenic portion
 CC of a prostate tumour protein. The encoded immunogen, or the DNA itself,
 CC can be used as a vaccine for the treatment of prostate cancer. The DNA
 CC was identified by analysis of a subcloned cDNA library obtained by
 CC subtracting a prostate tumour cDNA expression library with a normal
 CC tissue cDNA library.
 SO Sequence 852 BP; 174 A; 195 C; 206 G; 274 T;

Query Match 3.2%; Score 60.6; DB 1; Length 852;
 Best Local Similarity 56.2%; Pred. No. 0.00043;

Matches 114; Conservative 0; Mismatches 89; Indels 0; Gaps 0;

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QY 1163 caccctggcagcttccctcaagaggtgtctactcaagaagcccccacagctatgtccgctcatc 1222
DB 368 CACTATAGGAAATTCCTCCGACATCCCTCTTGACCCACAGACACCAAGCCAGATTAATA 329
QY 1223 aggtgtgtcaacaagaagagagctcttccaaattgagagctcagccaggtgcccgg 1282
DB 328 AATGCGAAGACGCACTGAGGCGCTCTTCAGTGTCTTGAAATCAGAGCAATGCTGTAC 269
QY 1283 ctgtgtgggacatcccaagaacgcgtccgcagcttctcagacagcgcgcgtccatc 1342
DB 268 CTATGGGGTAAAGAAAGAACACAGACGACATGACCTATGAAAGACCGCAGCTATC 209
QY 1343 ccgcagctatctcaagagagcatt 1365
DB 208 AGATTAATCTCAAAAGGAAT 186

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RESULT 14

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V1532915530 standard: cDNA; 2975 BP.
AC V15530:
DT 22-MAY-1998 (first entry)
DE Human lymphoid-specific transcription factor NERF-1 cDNA.
KW ets-related; human; lymphoid-specific transcription factor; NERF-1;
OS screening assay: ds.
FH Key 1 Location/Qualifiers
FT CDS
FT ID: 106
FT /tag= a
FT /product= NERF-1
PN US9721113.A.
PD 24-FEB-1998. 368281.
PR 03-JAN-1995; US-368281.
PR (HUMAN) HUMAN GENOME SCI INC.
PR KUNSCHE CA, LIBERMAN TA, OELTGEN JP.
DR P-PSDB: W47237.
DR P-PSDB: W47237.
PT NERF-1 and NERF-2 nucleic acids - encode ets-related human
PT lymphoid-specific transcription factors
PS Claim 1: Columns 17-22; 17pp: English.
CC The present sequence encodes the ets-related lymphoid-specific
CC transcription factor NERF-1, which can be used in screening assays
CC for drug that modulate NERF activity, and to treat patients having
CC need of NERF-1 ets-related factors are involved in, e.g. cancer
CC development, retrovirus replication, T-cell cytokine production,
CC cell cycle effects, growth regulation and cell differentiation.
SQ Sequence 2975 BP; 958 A; 575 C; 614 G; 828 T;

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Query Match 3.2%; Score 60.4; DB 1; Length 2975;

Best Local Similarity 53.9%; Pred. No. 0.0056;

Matches 124; Conservative 0; Mismatches 106; Indels 0; Gaps 0;

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QY 1151 gggagagccatccacctgtgagcttccaaaggtgtctccaaagcccaagcgtat 1210
DB 518 GGAACACACACCTATTGTGGAGTTCCTTTGATCTACTCTCAAGTAAATAACTGTG 577
QY 1211 ggcgcctcatatgtgtgtcacaagaagagagcattcaaatgtgagaccctcc 1270
DB 578 CCCAGGATATATTAAGTACACAGAGAAAGGATATATCAAGCTGATCAAAAG 637
QY 1271 caggtgtgcccgtgtgtggagctcgcagaagacgctccgcctgaactgaagcagcg 1330
DB 638 GCTGTCTTAAGCTTTGGGAAAGCATTAAGAACACACACATGATGAAACATG 697
QY 1331 agccgcctccatccgcagctattacaagaagagcattcgcagagccaga 1380
DB 698 GGACGAGCTTGGATCTACTACCAAGGCAATTTCTGCAAAAGTTGA 747

```

RESULT 15

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V15530
ID V15530 standard: cDNA; 3240 BP.
AC V15530:
DT 22-MAY-1998 (first entry)
DE Human lymphoid-specific transcription factor NERF-2 cDNA.
KW ets-related; human; lymphoid-specific transcription factor; NERF-2;
OS screening assay: ds.
FH Key 1 Location/Qualifiers
FT CDS
FT ID: 1952
FT /tag= a
FT /product= NERF-2
PN US9721113.A.
PD 24-FEB-1998. 368281.
PR 03-JAN-1995; US-368281.
PR (HUMAN) HUMAN GENOME SCI INC.
PR KUNSCHE CA, LIBERMAN TA, OELTGEN JP.
DR P-PSDB: W47238.
DR P-PSDB: W47238.
PT NERF-1 and NERF-2 nucleic acids - encode ets-related human
PT lymphoid-specific transcription factors
PS Claim 2: Columns 23-28; 17pp: English.
CC The present sequence encodes the ets-related lymphoid-specific
CC transcription factor NERF-2, which can be used in screening assays
CC for drug that modulate NERF activity, and to treat patients having
CC need of NERF-2 ets-related factors are involved in, e.g. cancer
CC development, retrovirus replication, T-cell cytokine production,
CC cell cycle effects, growth regulation and cell differentiation.
SQ Sequence 3240 BP; 1070 A; 592 C; 662 G; 916 T;

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Query Match 3.2%; Score 60.4; DB 1; Length 3240;

Best Local Similarity 53.9%; Pred. No. 0.0057;

Matches 124; Conservative 0; Mismatches 106; Indels 0; Gaps 0;

```

QY 1151 gggagagccatccacctgtgagcttccaaaggtgtctccaaagcccaagcgtat 1210
DB 783 GGAACACACACCTATTGTGGAGTTCCTTTGATCTACTCTCAAGTAAATAACTGTG 842
QY 1211 ggcgcctcatatgtgtgtcacaagaagagagcattcaaatgtgagaccctcc 1270
DB 843 CCCAGGATATATTAAGTACACAGAGAAAGGATATATCAAGCTGATCAAAAG 902
QY 1271 caggtgtgcccgtgtgtggagctcgcagaagacgctccgcctgaactgaagcagcg 1330
DB 903 GCTGTCTTAAGCTTTGGGAAAGCATTAAGAACACACACATGATGAAACATG 962
QY 1331 agccgcctccatccgcagctattacaagaagagcattcgcagagccaga 1380
DB 963 GGACGAGCTTGGATCTACTACCAAGGCAATTTCTGCAAAAGTTGA 1012

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Job time: 1640 sec